## **Amendments to the Claims**

This claim listing will replace all prior versions of claims and claim listings in the application:

## WHAT IS CLAIMED:

1. (Currently amended) A compound structurally represented by Formula I

$$R^{1}$$

Formula (I)

or pharmaceutically acceptable salts thereof wherein:

R<sup>1</sup> is

-CH<sub>2</sub>N  $R^3R^4$ ,

-CONR<sup>3</sup>R<sup>4</sup>,

 $R^2$  is

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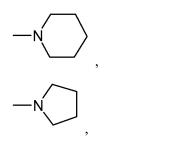
-Hydrogen,

 $-NH-(C_1-C_6)$  alkyl,

-NH-(C<sub>1</sub>-C<sub>4</sub>) alkylene-phenyl,

-NH(C3-C6)cycloakyl,

 $-NR^3R^4$ ,



wherein;

R<sup>3</sup> is hydrogen,

 $-(C_1-C_4)$  alkyl,

R<sup>4</sup> is

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 $-(C_1-C_4)$  alkyl,

-(C<sub>1</sub>-C<sub>4</sub>) alkylene -phenyl,

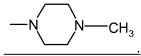
wherein  $R^3$  and  $R^4$  can cyclize to form, together with the nitrogen to which they are attached, a five or six-membered ring, wherein optionally one of the carbons of the ring formed by said nitrogen,  $R^3$ , and  $R^4$ , is replaced by a nitrogen or oxygen, and wherein said ring is optionally further substituted by  $R^5$ , and  $R^5$  is hydrogen,

-( $C_1$ - $C_4$ ) alkyl, wherein optionally  $R^5$  forms a 3 to five membered ring with the nitrogen containing ring to which it is attached,

-(C<sub>1</sub>-C<sub>4</sub>) alkylene –N-pyrrolidinyl,

-(C<sub>1</sub>-C<sub>4</sub>) alkylene –N- piperidinyl;

provided that when R1 is -CH2N R3R4 or -CONR3R4, then R2 is



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2. (Original) A compound structurally represented by Formula II,

or pharmaceutically acceptable salts thereof wherein:

Z is -carbonyl-, or -CH<sub>2</sub>-,

 $R^2$  is

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-Hydrogen,

 $-NH-(C_1-C_6)$  alkyl,

-NH-(C<sub>1</sub>-C<sub>4</sub>) alkylene-phenyl,

-NH(C<sub>3</sub>-C<sub>6</sub>)cycloakyl,

 $-NR^3R^4$ 

$$-N$$
 $N$ 
 $CH_3$ 

$$-N$$

$$-N$$

wherein;

15 R<sup>3</sup> is hydrogen,

$$-(C_1-C_4)$$
 alkyl,

 $R^4$  is

-(C<sub>1</sub>-C<sub>4</sub>) alkylene -phenyl,

wherein R<sup>3</sup> and R<sup>4</sup> can cyclize to form, together with the nitrogen to which they are attached, a five or six-membered ring, wherein optionally one of the carbons of the ring formed by said nitrogen, R<sup>3</sup>, and R<sup>4</sup>, is replaced by a nitrogen or oxygen, and wherein said ring is optionally further substituted by R<sup>5</sup>, and

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R<sup>5</sup> is hydrogen,

-( $C_1$ - $C_4$ ) alkyl, wherein optionally  $R^5$  forms a 3 to five membered ring with the nitrogen containing ring to which it is attached,

-(C<sub>1</sub>-C<sub>4</sub>) alkylene –N-pyrrolidinyl,

-(C<sub>1</sub>-C<sub>4</sub>) alkylene –N- piperidinyl.

- 3. (Original) The compound of claim 1, wherein R<sup>1</sup> is CONR<sup>3</sup>R<sup>4</sup>, and R<sup>3</sup> and R<sup>4</sup> cyclize to form, together with the nitrogen to which they are attached, a five membered ring, and said ring is further substituted by -CH<sub>2</sub>- pyrrolidinyl.
- 4. (Original) The compound of claim 1, wherein R<sup>1</sup> is CH<sub>2</sub>NR<sup>3</sup>R<sup>4</sup>, and R<sup>3</sup> and R<sup>4</sup> cyclize to form, together with the nitrogen to which they are attached, a five membered ring, and said ring is further substituted by -CH<sub>2</sub>- pyrrolidinyl.
- 5. (Original) The compound of claim 3 wherein R<sup>2</sup> is N R<sup>3</sup>R<sup>4</sup>, and R<sup>3</sup> and R<sup>4</sup> cyclize to form, together with the nitrogen to which they are attached, a five membered ring.
- 6. (Original) The compound of claim 4 wherein R<sup>2</sup> is N R<sup>3</sup>R<sup>4</sup>, and R<sup>3</sup> and R<sup>4</sup> cyclize to form, together with the nitrogen to which they are attached, a five membered ring.
  - 7. (Currently amended) The compound of claim 1, further represented by any one of the formula (Example 1) to (Example 18) selected from the group consisting of:

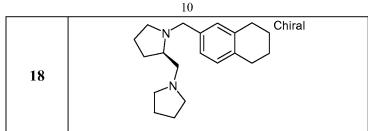
Example Number

1

	7
2	O Chiral
3	Chiral N
4	Chiral
5	Chiral
6	Chiral

	8
7	P F F
8	
9	Chiral
10	
11	

	9
12	
13	Chiral
14	Chiral
15	
16	Chiral
17	Chiral



or a pharmaceutically acceptable salt or solvate thereof.

- 8. (Currently amended) A pharmaceutical composition which comprises a compound of any of claims 1–7 and a pharmaceutically acceptable carrier.
- 9. (Currently amended) A method for treatment or prevention of a cognitive disorder 5 which comprises administering to a subject in need of such treatment or prevention an effective amount of a compound of any of Claims 1-7.
  - 10. (Original) The method of Claim 9 wherein the antagonist is a pharmaceutical composition of claim 8.
- 11. (Currently amended) A method for treatment or prevention of obesity which 10 comprises administering to a subject in need of such treatment or prevention an effective amount of a compound of any of Claims 1—7.
  - 12. (Original) The method of Claim 11 wherein the antagonist is a pharmaceutical composition of claim 8.
- 13. (Currently amended) A method for treatment or prevention of a wakefulness 15 disorder which comprises administering to a subject in need of such treatment or prevention an effective amount of a compound of any of Claims 1—7.
  - 14. (Original) The method of Claim 13 wherein the antagonist is a pharmaceutical composition of claim 8.
- 20 15. (Canceled).
  - 16. (New) A compound selected from the group consisting of:

	11
Example	
Number	
7	N P F F F
10	
11	
12	

or a pharmaceutically acceptable salt thereof.